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APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.
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10/824,593

04/15/2004

Raymond Pratt

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6645

26694

7590

07/15/2008

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EXAMINER

CHANNAVAJJALA, LAKSHMI SARADA

ART UNIT

PAPER NUMBER

1611

MAIL DATE

DELIVERY MODE

07/15/2008

PAPER

**Please find below and/or attached an Office communication concerning this application or proceeding.**

The time period for reply, if any, is set in the attached communication.

<b>Office Action Summary</b>	<b>Application No.</b> 10/824,593	<b>Applicant(s)</b> PRATT ET AL.	
	<b>Examiner</b> Lakshmi S. Channavajjala	<b>Art Unit</b> 1611	

-- The MAILING DATE of this communication appears on the cover sheet with the correspondence address --

### Period for Reply

A SHORTENED STATUTORY PERIOD FOR REPLY IS SET TO EXPIRE 3 MONTH(S) OR THIRTY (30) DAYS, WHICHEVER IS LONGER, FROM THE MAILING DATE OF THIS COMMUNICATION.

- Extensions of time may be available under the provisions of 37 CFR 1.136(a). In no event, however, may a reply be timely filed after SIX (6) MONTHS from the mailing date of this communication.
- If NO period for reply is specified above, the maximum statutory period will apply and will expire SIX (6) MONTHS from the mailing date of this communication.
- Failure to reply within the set or extended period for reply will, by statute, cause the application to become ABANDONED (35 U.S.C. § 133). Any reply received by the Office later than three months after the mailing date of this communication, even if timely filed, may reduce any earned patent term adjustment. See 37 CFR 1.704(b).

### Status

- 1) ☒ Responsive to communication(s) filed on 26 March 2008.
- 2a) ☐ This action is **FINAL**.                      2b) ☒ This action is non-final.
- 3) ☐ Since this application is in condition for allowance except for formal matters, prosecution as to the merits is closed in accordance with the practice under *Ex parte Quayle*, 1935 C.D. 11, 453 O.G. 213.

### Disposition of Claims

- 4) ☒ Claim(s) 25-45 is/are pending in the application.
- 4a) Of the above claim(s) \_\_\_\_\_ is/are withdrawn from consideration.
- 5) ☐ Claim(s) \_\_\_\_\_ is/are allowed.
- 6) ☒ Claim(s) 25-45 is/are rejected.
- 7) ☐ Claim(s) \_\_\_\_\_ is/are objected to.
- 8) ☐ Claim(s) \_\_\_\_\_ are subject to restriction and/or election requirement.

### Application Papers

- 9) ☐ The specification is objected to by the Examiner.
- 10) ☐ The drawing(s) filed on \_\_\_\_\_ is/are: a) ☐ accepted or b) ☐ objected to by the Examiner.  
Applicant may not request that any objection to the drawing(s) be held in abeyance. See 37 CFR 1.85(a).  
Replacement drawing sheet(s) including the correction is required if the drawing(s) is objected to. See 37 CFR 1.121(d).
- 11) ☐ The oath or declaration is objected to by the Examiner. Note the attached Office Action or form PTO-152.

### Priority under 35 U.S.C. § 119

- 12) ☐ Acknowledgment is made of a claim for foreign priority under 35 U.S.C. § 119(a)-(d) or (f).
- a) ☐ All    b) ☐ Some \*    c) ☐ None of:
1. ☐ Certified copies of the priority documents have been received.
  2. ☐ Certified copies of the priority documents have been received in Application No. \_\_\_\_\_.
  3. ☐ Copies of the certified copies of the priority documents have been received in this National Stage application from the International Bureau (PCT Rule 17.2(a)).

\* See the attached detailed Office action for a list of the certified copies not received.

### Attachment(s)

- |  |   |
|--|---|
| 1) <input checked="" type="checkbox"/> Notice of References Cited (PTO-892)            | 4) <input type="checkbox"/> Interview Summary (PTO-413)           |
| 2) <input type="checkbox"/> Notice of Draftsperson's Patent Drawing Review (PTO-948)   | Paper No(s)/Mail Date. _____                                      |
| 3) <input checked="" type="checkbox"/> Information Disclosure Statement(s) (PTO/SB/08) | 5) <input type="checkbox"/> Notice of Informal Patent Application |
| Paper No(s)/Mail Date ( <u>1 page</u> ) <u>2-12-08</u> .                               | 6) <input type="checkbox"/> Other: _____                          |

### **DETAILED ACTION**

Receipt of RCE dated 3-26-08 is acknowledged.

#### ***Continued Examination Under 37 CFR 1.114***

1. A request for continued examination under 37 CFR 1.114, including the fee set forth in 37 CFR 1.17(e), was filed in this application after final rejection. Since this application is eligible for continued examination under 37 CFR 1.114, and the fee set forth in 37 CFR 1.17(e) has been timely paid, the finality of the previous Office action has been withdrawn pursuant to 37 CFR 1.114. Applicant's submission filed on 3-26-08 has been entered.

Claims 25-45 are pending in the instant application. Instant independent claims have been amended recite the drugs or substances that induce abuse.

#### ***Response to Amendment***

2. In response to applicants' amendment, the following new rejection has been applied. Applicants' arguments of 9-28-07 have been addressed in the action dated 12-17-07. Neither of the amendment of 12-12-07 nor the RCE of 3-26-08 is associated with arguments or remarks.

#### ***Claim Rejections - 35 USC § 103***

The text of those sections of Title 35, U.S. Code not included in this action can be found in a prior Office action.

Claims 25-45 are rejected less than 35 U.S.C. 103(a) as being unpatentable over WO 98/39000 (WO) in view of US 5,278,176 to Lin and Nath et al or over WO in view of Nath et al

WO teaches methods of treating disorders of attention or improving attention by administering an effective amount of a cholinesterase inhibitor. WO teaches that acetylcholine is a neural transmitter for transmitting messages across the synapse to a cholinergic message by stimulating the cholinergic receptor for neuronal messages such as memory (page 3, L 23-29). WO teaches that cholinesterase rapidly destroys acetylcholine resulting in a weak cholinergic stimulation, experienced as a memory loss, and states that one way to overcome the above loss is to interfere with the ability of cholinesterase to degrade acetylcholine, as by treatment with cholinesterase inhibitors (page 4, l 5-12). WO teaches all of the instant claimed compounds, including donepezil, for their cholinesterase inhibiting activity and thus inhibiting memory loss (pages 5+). WO fails to teach the claimed method of treating substance abuse or treating withdrawal symptoms or decreasing the rate of relapse.

Lin teaches selective and potent nicotinic agonists that are useful in treating dementias, attention disorders, and anxiety associated with cognitive impairment or substance abuse withdrawal characterized by decreased cholinergic function (abstract). Lin teaches that chronic alcoholism (reads on instant substance abuse, see instant claim 25) and the resultant brain disease such as Alzheimer's disease, is characterized by diffuse reductions in cortical cerebral blood flow in those brain regions where cholinergic neurons arise (col. 2, L 61-65). Lin further states that nicotine withdrawal

syndrome associated with tobacco use is characterized by craving for nicotine, irritability, frustration, anger, difficulty in concentration etc (col. 4, L 60-67). Lin suggests that symptoms associated with withdrawal of nicotine or compounds that act as nicotine agonists for acetylcholine receptors can alleviate other addictive substances.

Kish (abstract only) teaches Cognitive impairment has been reported in some chronic users of psychostimulants, raising the possibility that long-term drug exposure might damage brain neuronal systems, including the cholinergic system which is responsible for normal cognition. Kish reports that a measurement of the activity of choline acetyltransferase (ChAT), the marker enzyme for cholinergic neurons, in autopsied brain of chronic users of cocaine, methamphetamine, and, for comparison, heroin showed that as compared with the controls, mean ChAT levels were normal in all cortical and sub cortical brain areas examined. However, the two of 12 methamphetamine users, who had the highest brain/blood drug levels at autopsy, had a severe (up to 94%) depletion of ChAT activity in cerebral cortex, striatum, and thalamus. Based on the subjects examined in the present study, our neurochemical data, Kish states that the brain cholinergic neurone damage is unlikely to be a typical feature of chronic use of cocaine, methamphetamine, or heroin, but that exposure to very high doses of methamphetamine could impair, at least acutely, cognitive function requiring a normal nucleus basalis cholinergic neuronal system. Reduced brain ChAT might be explained in part by a hyperthermia-related mechanism as low ChAT levels have also been observed in brain of some patients with neuroleptic drug-associated hyperthermia.

Studies of cognitive and brain cholinergic status in high dose users of MA are warranted. Kish does not teach donepezil for cognitive impairment or cholinergic status.

It would have been obvious for one of an ordinary skill in the art at the time of the instant invention to use the cholinesterase inhibitors such as donepezil and other compounds of treating substance abuse or withdrawal symptoms associated with substance abuse because both Lin and Kish emphasizes that the cholinergic system plays a profound role in substance abuse or drug withdrawal symptoms that are associated with substances such as alcohol, cocaine or methamphetamine (which are claimed to cause substance abuse) and Lin states that cholinesterase inhibitors effectively interfere with and alleviate the substance abuse and the associated withdrawal symptoms. Even though Kish states that high doses of drugs impair cognitive functioning requiring a normal cholinergic neuronal system, instant claims are silent regarding the amount of substances that cause the withdrawal symptoms or symptoms of substance abuse. Hence, one of an ordinary skill in the art would have expected the cholinesterase inhibitors of WO to be effective in reducing the substance abuse or in interfering with the drug withdrawal symptoms caused by abusive drugs such as nicotine, cocaine, methamphetamine, alcohol etc.

While Lin does not teach all of the claimed drugs that result in the abuse or cause the withdrawal symptoms, Lin teaches the underlying mechanism that results in the claimed abuse and its withdrawal symptoms and hence a skilled artisan would have expected to effectively reduce substance abuse, caused by any substance, with the

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cholinesterase inhibitors of WO. Further, optimizing the amount and the route of administration of the compounds of WO, so as to reduce substance addiction and also prevent the return to drug seeking behavior would have been within the scope of a skilled artisan.

Any inquiry concerning this communication or earlier communications from the examiner should be directed to Lakshmi S. Channavajjala whose telephone number is 571-272-0591. The examiner can normally be reached on 9.00 AM -5.30 PM.

If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, Michael Woodward can be reached on 571-272-8373. The fax phone number for the organization where this application or proceeding is assigned is 571-273-8300.

Information regarding the status of an application may be obtained from the Patent Application Information Retrieval (PAIR) system. Status information for published applications may be obtained from either Private PAIR or Public PAIR. Status information for unpublished applications is available through Private PAIR only. For more information about the PAIR system, see <http://pair-direct.uspto.gov>. Should you have questions on access to the Private PAIR system, contact the Electronic Business Center (EBC) at 866-217-9197 (toll-free). If you would like assistance from a USPTO Customer Service Representative or access to the automated information system, call 800-786-9199 (IN USA OR CANADA) or 571-272-1000.

/Lakshmi S Channavajjala/  
Primary Examiner,  
Art Unit 1611  
June 23, 2008